SOURCE CODE

```
/* procedure:
               dsm_rindex.sas revision:
/* authored_by: doug mcnair modified_by:
/* created_date: 04-JAN-2003 modified_date:
data transform for adjusting health service
                 for variations in rurality distance or access */
/* environment: H:\cerdsm\NHS\test
/* Read in data file. Assign the name "pop" to catchment pop...
/* Assign the name "d nhs" to distance to care category to be
/* distance-index adjusted.
/* Only the infile statement and the the input statement need to */
/* be modified.
/* This is a locally normal index and assigns a quantitative
/* measure of rurality to each participant in a study. Data req'd */
/* to compute the index are minimal; only two variables need be: */
/* measured. Analysis effort required to compute the index is */
/* negligible, and the tradeoff between data collection and data */
/* analysis is defensible. Data collection is generally more */
/* expensive than data analysis so minimize that. The two required*/
/* variables, county population and distance to care location, are*/
/* optimally transformed to achieve normality and weighted for */
/* validity. Measure of departure from normality is automatically */
/* obtained while constructing the index. Positive score reflects */
/* rural loc or urban loc with compromised access. Negative score */
/* reflects an urban residence with good access relative to the . */
/* group under study. A score of zero reflects average distance
/* or accessibility of health services, rural or otherwise.
/* Reliability and construct validity were examined using two:
/* data sets.
/*
/* The choice of a distance measure to gauge the distance between */
/* two distributions, in this case Gaussians, is critical.*/
/* The Kullback-Leibler, Bhattacharyya and arc-cosine distances */
/* are all related to maximum likelihood, and reduce to the */
/* Mahalanobis distance when used to measure the distance between */
/* two Gaussian distributions with equal covariance matrices.
/* The advantage of the arc cosine distance is that it has a
/* closed form expression, implemented in IML.
data distance;
    infile 'c:\0_cerdsm\health_econ\rurality\cancerpop.dat';
    input case id d nhs pop;
    retain d nhs pop;
    use=1;
    if d nhs = 0 then d nhs=.5;
    if d nhs =. then use=0;
    if pop = . then use=0;
    keep d_nhs pop use;
```

```
if use = 1 then output;
 proc means;
 proc freq;
     tables d nhs;
 proc iml;
    use distance; reset noname; read all var _num into YY;
    step=.05; mino=step; maxo=4.; guess={1, 1};
    k=2; c=\{.5, -1\}; n=nrow(YY);
    do i=1 to k; YY[,i]=YY[,i]/max(YY[,i]); end;
    j=(1:n)'; p=(j-.5)/n; pi=arcos(-1);
    X=J(n,k,0); X1=J(n,k,0); X2=J(n,k,0);
    eps=1/2**26;
    start deriv;
    tau=lam-1; omega=(ssq(tau)); romega=sqrt(omega); IMP=I(k);
    if romega >= eps then do;
    tau=tau/romega; IMP=IMP-tau*tau'; end;
    do j=1 to k;
      y=YY[,j];
      if lam[j] =0 then do;
        t=log(y);
        t1=t#t/2;
        t2=2*t1#t/3;
                      end;
      else do;
        t=sign(lam[j])*y##lam[j];
        t1=t\#\log(y); t2=t1\#\log(y); end;
      t=t-J(n,1,sum(t)/n); v0=ssq(t); xj=t/sqrt(v0); X[,j]=xj;
      t1=t1-J(n,1,sum(t1)/n); t2=t2-J(n,1,sum(t2)/n);
      v01=xj'*t1; v02=xj'*t2;
      x1j=(t1-xj*v01)/sqrt(v0); X1[,j]=x1j; v11=x1j'*t1;
      x2j=(t2-xj*v02-2*x1j*v01-xj*v11)/sqrt(v0); X2[,j]=x2j;
      end;
      z=X*c; s2=z*z/(n-1); s=sqrt(s2); z=z/s;
      ii=rank(z); B=z; z[ii]=B;
      B=X; X[ii,]=B; B=X1; X1[ii,]=B; B=X2; X2[ii,]=B;
      G=sqrt(omega/s2)*diag(c)*(X1'-(X1'*z)*z'/(n-1));
      Phi=probnorm(z);
      Phi = Phi\#(eps <= Phi)\#(Phi <= 1-eps) + J(n,1,eps)\#(Phi < eps)
          + J(n,1,1-eps) # (1-eps < Phi);
      logphi=log(Phi); logphi1=log(1-Phi);
      ph=exp(-z#z/2)/sqrt(2*pi); logph=-(z#z+J(n,1,log(2*pi)))/2;
      A2n=2*p'*logphi1-2*p'*logphi-2*sum(logphi1)-n;
      a1=-2*exp(logph-logphi-logphi1)#(p-Phi);
      a2=2*exp(2*logph-2*logphi-2*logphi1)#((p-Phi)#(p-Phi)+p#(1-p));
      a2=a2+2*exp(logph-logphi-logphi1)#z#(p-Phi);
      d1=G*a1; f=IMP*d1;
      d2=(omega/s)*diag(c)*(X2`*a1-(X2`*z)*(z`*a1)/(n-1));
      d3=sqrt (omega/s2) *IMP*diag(c) *X2 \*z;
      H=-(f*tau'+tau*f'); H1=diag(d2)-(tau'*d1)*I(k)-(d1*d3'+d3*d1')/(n-d1*d3'+d3*d1')
1);
      H1=H1-G*G**(z**a1)/(n-1)+(G#(J(k,1,1)*a2*))*G*;
      H=H+IMP*H1*IMP:
      finish;
     start optimize;
```

```
delta=1; A2n0=n;
 do while (delta > 1.e-5);
 run deriv;
 call gsorth(T,u,lindep,I(2)-tau*tau'); T=T[,1];
 b=-inv(T'*H*T)*(T'*f);
 tau=tau+T*b;
 tau=tau/sqrt(ssq(tau));
 lam=1+tau*romega;
 delta=n*abs(A2n0-A2n);
 A2n0=A2n;
 end;
 finish;
start search;
filename out 'power.out';
file out;
lam={1,1}; run deriv;
put @2 romega 7.4 +3 A2n 9.4+3 (lam[1]) 7.4+3 (lam[2]) 7.4+3;
lam=guess-.001;
do romega=mino to maxo by step;
omega=romega**2;
tau=lam-1; tau=tau/sqrt(ssq(tau)); lam=1+romega*tau;
run optimize;
put @2 romega 7.4 +3 A2n 9.4+3 (lam[1]) 7.4+3 (lam[2]) 7.4+3;
end;
closefile out;
finish;
lam={0.2,-.1}; run deriv; print lam A2n;
lam={0,-.15}; run deriv; print lam A2n;
lam={0,-.2}; run deriv; print lam A2n;
lam={1.,1.}; run deriv; print lam A2n;
lam={0.,0.}; run deriv; print lam A2n;
run search;
```

SOURCE CODE

```
/* purpose: calculate risk-adjusted area rates for indicators*/
            variations in rurality distance or access */
/*
              adj for age and dcat
                                                */
/* environment: H:\cerdsm\NHS\test
                                                */
filename control 'h:\cerdsm\nhs\control.sas';
%include control;
%macro area;
 %if &msalevl eq 1 %then msa;
    %else hospstco;
%mend;
%macro cty2msa;
  %if &msalevl eq 1 %then %do;
    label msa ='msa %area assigned from hospstco';
    msa = input( put(hospstco, area.), 5.);
  %end;
%mend;
%macro sum2msa;
  %if &msalevl eq 1 %then %do;
     proc summary nway;
        class msa popcat dcat;
        var pop;
        output out=qipop (drop=_type__freq ) sum=;
  %end;
%mend;
                ------
                add pop denominator
       ------
proc
     sort data=in3.&infilea3. (keep=%area) out=%area nodupkey;
by
     %area;
run;
data
     qipop;
length hospstco 8 popcat 3 dcat $1;
infile popfile3 missover;
input hospstco popcat dcat pop;
%cty2msa
run:
%sum2msa
```

```
proc sort data=qipop;
 by
      %area popcat dcat;
 run;
data
      qipop;
merge %area(in=x) qipop;
by
      %area;
if x;
run;
 /* -----*/
 /* select obs for each indicator from previous step in turn.
 /* each iteration passes vars that control covars for that indic: */
      n - obs number from covar files
                                                       */
         also identifies format used to index covars
                                                      . */
      pq - indicator name without prefix
/*
      a - number of demographic categories (age and d nhs)
      s - determines whether age*d interactions are needed
                                                    * /
      o - determines omitted age cat in format
   %macro mod3(n,pq,a,s,o);
/* -----*/
/* SET creates TEMP1, containing the dep var TPQ and indep vars */
/* used in the regression. append to enctr data one obs per %area */
/* and demographic grp
      temp1(keep=key %area t&pq. d agel-age%eval(2*&a.) aidx pop
mergevar);
length age1-age%eval(2*&a.) aidx 8;
set in3.&infilea3.(keep=key %area t&pq. dcat agecat age)
      qipop;
/* -----*/
/* retain only enctr/area/demographic grp needed for this indicator*/
/* -----*/
n = \&n.;
if %area eq . then delete;
if pop > 0 and n notin (2,9) then t&pq. = 0;
if agecat in ('1') or popcat in (1,2,3,4) then do;
  if n in (1,3,5,7,8,13,14,15,16)
   then t&pq. = .;
end;
else if agecat in ('2') or popcat in (5,6,7,8) then do;
  if n in (4,6,9)
   then t&pq. = .;
end;
else if agecat in ('3','4') or
      popcat in (9,10,11,12,13,14,15,16,17,18) then do;
  if n in (4,6,9)
   then t&pq. = .;
```

```
end;
if t&pq. ne .;
/* initialize covar dummies to zero
/* ------
array agex{%eval(2*&a.)} age1-age%eval(2*&a.);
d = 0;
do i = 1 to eval(2*&a.);
 agex(i)=0;
end;
                     ----*/
/* set index for age array (AIDX). categories vary by indicator and*/
/* are determined by format corresponding to N
/* ------
if age > 0 then aidx = put(age,a&n.fmt.);
else if pop > 0 then do;
 aidx = popcat;
 if aidx = \&o. then aidx = 0;
else aidx = 0;
/* -----*/
/* set appropriate covariate dummy to 1 for this enctr */
/* -----*/
if dcat in (&s.) then d = 1;
if aidx > 0 then do; agex(aidx) = 1; agex(&a.+aidx) = 1 * d; end;
mergevar = 1;
run;
proc
    sort data=temp1;
by %area d aidx pop;
run;
/* -----*/
/* subtract number of encounters from %area/demographic group pop */
/* and set weight of 1 for each obs from enctr data, and a */
/* weight of (pop-encounters) for the appended obs for each group */
/* -----*/
data temp1(keep=key %area t&pq. d age1-age%eval(2*&a.) wht mergevar);
set temp1;
by %area d aidx;
retain wcnt 0;
if first.aidx then wcnt = 0;
if pop > 0 then wht = pop - wcnt;
else wht = 1;
wcnt + 1;
```

```
run;
/* read overall mean file for indicator; F variables are filler */
  *----*/
     temp1m(drop=n f1-f%eval(40-(2*&a.)));
length n f1-f%eval(40-(2*\&a.)) 3 md mage1-mage%eval(2*\&a.) 8;
infile meansa missover lrecl=1000;
input n md magel-mage%eval(2*&a.) f1-f%eval(40-(2*&a.));
mergevar = 1;
if n = &n:
run;
/* subtract overall mean of vars from the vars on each obs
/* so that adj area rate is the rate for the average person */
/* -----*/
data
     temp1(keep=key %area t&pq. d age1-age%eval(2*&a.) wht);
merge temp1 temp1m;
by
     mergevar;
array covar{%eval(1+(2*&a.))} d age1-age%eval(2*&a.);
array means{%eval(1+(2*&a.))} md mage1-mage%eval(2*&a.);
do i = 1 to eval(1+(2*&a.));
 covar(i) = covar(i) - means(i);
end;
run;
           read regression coeffs for each covariate */
     temp2(keep=intercep d age1-age%eval(2*&a.) _type_ _model
        depvar t&pq.);
length intercep d age1-age%eval(2*&a.) f1-f%eval(40-(2*&a.)) t&pq. 8
     _type_ _model_ _depvar_ $8;
infile covara missover lrecl=1000;
input n intercep d agel-age%eval(2*&a.) f1-f%eval(40-(2*&a.))
     _depvar_ _model_ _type_ _rmse_ t&pq.;
if n = &n.;
run;
                   -----*/
/* calculate predicted indicator for each obs using actual covars */
/* and the estimated coeffs, on de-meaned sample, where %area: */
/* effects are removed by subtracting the predicted val from the */.
/* actual on each obs, and calculating the mean difference for the */
/* %area. See Green WH. Econometric Analysis 2e. 1990 pp. 466-9. */
/* -----*/
```

```
proc
       score data=temp1 score=temp2 type=parms out=temp1y;
var
       d age1-age%eval(2*&a.);
run;
data
       temply;
set
       temply;
ehat = t&pq. - mhat;
one = 1;
run;
proc
       summary data=temply ;
class %area;
var
       ehat one;
output out=r&pq. mean(ehat)=r&pq. sum(one)=p&pq.;
run;
%mend;
%MOD3(1,APQ01,18,'F',6);
%MOD3(2,APQ02,4,'F',0);
%MOD3 (3, APQ03, 18, 'F', 10);
%MOD3 (4, APQ04, 18, 'F', 1);
%MOD3 (5, APQ05, 18, 'F', 12);
%MOD3(6,APQ06,18,'F',1);
%MOD3(7,APQ07,18,'F',10);
%MOD3(8,APQ08,18,'F',13);
%MOD3(9,APQ09,2,'F',0);
%MOD3 (10, APQ10, 18, 'F', 1);
%MOD3 (11, APQ11, 18, 'F', 5);
%MOD3 (12, APQ12, 18, 'F', 6);
%MOD3(13,APQ13,18,'F',11);
%MOD3 (14, APQ14, 18, 'F', 8);
%MOD3 (15, APQ15, 18, 'F', 7);
%MOD3 (16, APQ16, 18, 'F', 11);
   merge % area adjusted rates for indicators
data riskadj;
length %area RAPQ01-RAPQ16 PAPQ01-PAPQ16 8;
merge RAPQ01(KEEP=%AREA RAPQ01 PAPQ01)
      RAPQ02(KEEP=%AREA RAPQ02 PAPQ02)
      RAPQ03 (KEEP=%AREA RAPQ03 PAPQ03)
      RAPQ04 (KEEP=%AREA RAPQ04 PAPO04)
      RAPQ05 (KEEP=%AREA RAPQ05 PAPQ05)
      RAPQ06 (KEEP=%AREA RAPQ06 PAPQ06)
    RAPQ07(KEEP=%AREA RAPQ07 PAPQ07)
      RAPQ08 (KEEP=%AREA RAPQ08 PAPQ08)
     RAPQ09 (KEEP=%AREA RAPQ09 PAPQ09)
      RAPQ10 (KEEP=%AREA RAPQ10 PAPQ10)
      RAPQ11 (KEEP=%AREA RAPQ11 PAPQ11)
      RAPQ12 (KEEP=%AREA RAPQ12 PAPQ12)
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```
RAPQ13 (KEEP=%AREA RAPQ13 PAPQ13)
       RAPQ14 (KEEP=%AREA RAPQ14 PAPO14)
       RAPQ15 (KEEP=%AREA RAPQ15 PAPQ15)
       RAPQ16 (KEEP=%AREA RAPQ16 PAPQ16);
by
       %area;
label
RAPQ01 = 'DIABETES SHORT TRM COMPLICATN (Risk Adj)'
RAPQ02 = 'PERFORATED APPENDIX
                                       (Risk Adj)'
RAPQ03 = 'DIABETES LONG TERM COMPLICATN (Risk Adj)'
RAPQ04 = 'PEDIATRIC ASTHMA
                                       (Risk Adj)'
RAPQ05 = 'COPD
                                       (Risk Adj)'
RAPQ06 = 'PEDIATRIC GASTROENTERITIS
                                       (Risk Adj)'
RAPQ07 = 'HYPERTENSION
                                       (Risk Adj)'
RAPQ08 = 'CONGESTIVE HEART FAILURE
                                       (Risk Adj)'
RAPQ09 = 'LOW BIRTH WEIGHT
                                       (Risk Adj)'
RAPO10 = 'DEHYDRATION
                                       (Risk Adj)'
RAPQ11 = 'BACTERIAL PNEUMONIA
                                       (Risk Adj)'
RAPQ12 = 'URINARY INFECTION
                                       (Risk Adj)'
RAPQ13 = 'NEW CANCER MGT DELAY>14D S/P
                                       (Risk Adj)'
RAPQ14 = 'DIABETES UNCONTROLLED
                                       (Risk Adj)'
RAPQ15 = 'ADULT ASTHMA
                                       (Risk Adj)'
RAPQ16 = 'LOWER EXTREMITY AMPUTATION
                                       (Risk Adj)'
label
PAPQ01 = 'DIABETES SHORT TRM CMPLICTN (Denominator)'
PAPQ02 = 'PERFORATED APPENDIX
                                     (Denominator)'
PAPQ03 = 'DIABETES LONG TRM CMPLICTN
                                     (Denominator)'
PAPQ04 = 'PEDIATRIC ASTHMA
                                     (Denominator)
PAPQ05 = 'COPD
                                     (Denominator)'
PAPQ06 = 'PEDIATRIC GASTROENTERITIS
                                     (Denominator)'
PAPQ07 = 'HYPERTENSION
                                     (Denominator)'
PAPQ08 = 'CONGESTIVE HEART FAILURE
                                     (Denominator)' -
PAPQ09 = 'LOW BIRTH WEIGHT
                                     (Denominator)'
PAPQ10 = 'DEHYDRATION
                                     (Denominator)
PAPQ11 = 'BACTERIAL PNEUMONIA
                                     (Denominator)'.
PAPQ12 = 'URINARY INFECTION
                                     (Denominator)
PAPQ13 = 'NEW CANCER MGT DELAY>14D S/P(Denominator)'
PAPQ14 = 'DIABETES UNCONTROLLED
                                     (Denominator)'
PAPQ15 = 'ADULT ASTHMA
                                     (Denominator)'.
PAPQ16 = 'LOWER EXTREMITY AMPUTATION (Denominator)'
array arry1{16} rapq01--rapq16;
do i = 1 to 16;
 if arry1(i) < 0 then arry1(i) = 0;
 else if arry1(i) > 1 then arry1(i) = 1;
end;
drop i;
run;
              ----*/
/* smoothed demeaned rates = area rate * (signal / signal + noise) */.
/* -----*/
```

```
data
        smoothed(DROP=RAPQ01-RAPQ16);
 set
       riskadj;
 %include msx;
 array arrya{16} RAPQ01-RAPQ16;
 array arryn{16} PAPQ01-PAPQ16;
array arryx{16} SAPQ01-SAPQ16;
do i = 1 to 16;
  arryx(i) = arrya(i) - arrya3(i);
  if arryn(i) ne 0 and arrya2(i) ne 0 then
     arryx(i) = arryx(i) * (arrya2(i) / (arrya2(i) + (arrya1(i)))
arryn(i))));
 else arryx(i)=.;
  arryx(i) = arryx(i) + arrya3(i);
label
SAPQ01 = 'DIABETES SHORT TRM COMPLICATN (Smoothed)'
SAPQ02 = 'PERFORATED APPENDIX
                                         (Smoothed)'
SAPQ03 = 'DIABETES LONG TERM COMPLICATN (Smoothed)'
SAPQ04 = 'PEDIATRIC ASTHMA
                                         (Smoothed)'
SAPQ05 = 'COPD
                                         (Smoothed)'
SAPQ06 = 'PEDIATRIC GASTROENTERITIS
                                         (Smoothed) '
SAPQ07 = 'HYPERTENSION
                                         (Smoothed)'
SAPQ08 = 'CONGESTIVE HEART FAILURE
                                         (Smoothed)'
SAPQ09 = 'LOW BIRTH WEIGHT
                                         (Smoothed)'
SAPQ10 = 'DEHYDRATION
                                         (Smoothed)'
SAPQ11 = 'BACTERIAL PNEUMONIA
                                         (Smoothed)'
SAPQ12 = 'URINARY INFECTION
                                         (Smoothed)'
SAPQ13 = 'NEW CANCER MGT DELAY>14D S/P
                                         (Smoothed)'
SAPQ14 = 'DIABETES UNCONTROLLED
                                         (Smoothed)'
SAPQ15 = 'ADULT ASTHMA
                                         (Smoothed)'
SAPQ16 = 'LOWER EXTREMITY AMPUTATION
                                         (Smoothed)'
drop i;
run;
                              merge rates
data
       out3.&outfila3.;
merge in3.&infilea2. (where=(_type_ in(0,8))
    keep=%area _type_ tapq01-tapq16 papq01-papq16 oapq01-oapq16)
         riskadj (keep=%area rapq01-rapq16)
         smoothed(keep=%area sapq01-sapq16) ;
by
       %area;
run;
                             output rates
```

CANCER POP. DATA

100	35	12083
101	20	11697
102	. 15	5046
103	30	59218
104	20	9505
105	25	113419
106	20	8669
107	90	12083
.108	10	1383
109	10	8669
110	5	14562
111	10	9505
112	45	8669
113	15	12083
114	25	8239
150	20	113419
151	45	59218
152	8	5452
153	45	1503
154	120	25010
155	15	11337
156	5	6536
157	5	9505
-158	.5	5163
159	25	8669
160	45	78687
161	30	6536
162	5	10505
163	15	4106
164	40	12083